



(19) Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) Publication number:

0 446 685 A1

(12)

EUROPEAN PATENT APPLICATION

(21) Application number: **91102550.0**

(51) Int. Cl. 5: **C08K 13/02, C08L 57/08,**
//(C08K13/02,3:26,5:07,5:09)

(22) Date of filing: **21.02.91**

(30) Priority: **15.03.90 JP 65154/90**

(43) Date of publication of application:
18.09.91 Bulletin 91/38

(84) Designated Contracting States:
BE CH DE FR GB LI NL

(71) Applicant: **ASAHI DENKA KOGYO KABUSHIKI KAISHA**
2-35, Higashiogu 7-chome
Arakawa-ku Tokyo 116(JP)

(72) Inventor: **Sugawara, Yuji**
8-7-7 Tsutsuo
Kuwana-shi, Mie-ken(JP)
Inventor: **Kobe, Shinji**
4-301, 6-1249-1 Mitsuhashi
Omiya-shi, Saitama-ken(JP)

(74) Representative: **Patentanwälte Grünecker,
Kinkeldey, Stockmair & Partner**
Maximilianstrasse 58
W-8000 München 22(DE)

(54) **Halogenated resin composition.**

(57) A halogenated resin composition whose heat stability, weatherability, initial coloration, thermal coloration and adhesiveness are improved by adding thereto an overbased alkaline earth metal carboxylate/carbonate together with a β -diketone compound or a metal salt thereof.

EP 0 446 685 A1

Background of the Invention:

[Field of the Invention]

5 This invention relates to a halogenated resin composition which is stabilized by adding thereto a specific stabilizer system. More particularly, it relates to a halogenated resin composition whose heat stability, weatherability, initial coloration, thermal coloration and adhesiveness are improved by adding thereto an overbased a alkaline earth metal carboxylate/carbonate together with a β -diketone compound or a metal salt thereof.

10

[Description of the Prior Art]

It is known that a halogenated resin, which is less stable to heat and light, would frequently undergo decomposition mainly caused by dehydrohalogenation in a thermal molding step or when used at a relatively high temperature or upon exposure to ultraviolet rays, for example, outdoor usage.

15 Thus attempts have been made to improve the stability of a halogenated resin by adding several heat stabilizers such as metal salts of organic acids, inorganic basic metal compounds, organotin compounds, organic phosphite compounds, epoxy compounds or β -diketone compounds thereto.

Among these heat stabilizers, metal salts of organic acids have been used as a fundamental stabilizer 20 for halogenated resins. Further, there has been already proposed the use of a metal salt of an organic acid, wherein a metal base is used in a stoichiometrically excessive amount (i.e., so-called "overbased complex"), as described in, for example, Japanese Patent Publication No. 49019/1977.

25 Although these stabilizers comprising metal salts of organic acids are effective in the improvement of the heat stability over a prolonged period of time, they are disadvantageous from the viewpoint of coloration, in particular, coloration in the initial stage of heating or after the completion of heating.

It is further known that β -diketone compounds are stabilizers which can mainly improve coloration. There has been proposed in, for example, Japanese Patent Publication No. 8786/1967, Japanese Patent Laid-Open No. 34939/1976, Japanese Patent Laid-Open No. 95447/1976 and Japanese Patent Laid-Open No. 9847/1978 the use of a β -diketone compound or a metal salt thereof and a metal soap.

30 Although these β -diketone compounds are effective in the improvement of coloration, they are less effective in the improvement of heat stability over a prolonged period of time.

Thus Japanese Patent Publication No. 6465/1982 has proposed to use a β -diketone compound or a metal complex salt thereof together with an overbased alkali metal or alkaline earth metal organic sulfonate or phenolate complex to thereby simultaneously improve the heat stability and the coloration.

35 Recently, the use of halogenated resins have been diversified and thus the processing conditions and the environment of use therefor have become more and more severe, which makes the stabilizing effect of known stabilizers insufficient. Further, there has been frequently observed the use of a composite material comprising a halogenated resin and a different kind of material, for example, a metal. However a conventional compounding system has only a limited adhesiveness to a different kind of material and, as a result, readily peels off, which results in an additional disadvantage.

40 Thus it has been urgently required to find out a stabilizer system for a halogenated resin whereby not only the heat stability at a processing temperature and coloration of the halogenated resin but also the heat stability at a relatively high temperature of approximately 100 °C (i.e., the so-called heat aging characteristics) and the adhesiveness to a different kind of material can be improved.

45

Summary of the Invention:

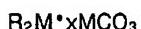
Under these circumstances, the present inventors have conducted extensive studies in order to find out a stabilizer whereby not only a halogenated resin can be stabilized over a long time even under severe 50 conditions of use but also the adhesiveness thereof to a different kind of material can be improved. As a result, they have found out that a halogenated resin composition excellent in heat stability, coloration and adhesiveness can be obtained by adding an overbased alkaline earth metal carboxylate/ carbonate and a β -diketone compound or a metal salt thereof to a halogenated resin.

Accordingly, the present invention provides a stabilized halogenated resin composition which comprises 55 100 parts by weight of a halogenated resin and, added thereto, (a) 0.05 to 10 parts by weight of at least one overbased alkaline earth metal carboxylate/carbonate and (b) 0.001 to 5 parts by weight of at least one β -diketone compound or a metal salt thereof.

Detailed Description of the Invention:

The overbased alkaline earth metal carboxylate/ carbonate to be used in the present invention is a compound which is schematically represented by the following formula:

5



wherein R represents an organic carboxylic acid residue;

M represents an alkaline earth metal; and

10 x represents the excessive amount of the metal and is a positive number exceeding 0.

Examples of the alkaline earth metal represented by M herein include barium, calcium, magnesium and strontium. x generally ranges from 0.1 to 60, preferably from 0.5 to 30, though it is not particularly restricted as long as it exceeds 0. Examples of the organic acid residue represented by R include residues of monocarboxylic acids such as caproic acid, caprylic acid, pelargonic acid, 2-ethylhexylic acid, capric acid, neodecanoic acid, undecylenic acid, lauric acid, myristic acid, palmitic acid, stearic acid, isostearic acid, 12-hydroxystearic acid, chlorostearic acid, 12-ketostearic acid, phenylstearic acid, ricinolic acid, linoleic acid, linolenic acid, oleic acid, arachic acid, behenic acid, erucic acid, brassidic acid and similar acids, mixtures of the above-cited acids with natural acids such as tallow fatty acids, coconut oil fatty acids, tung oil fatty acids, soybean oil fatty acids and cottonseed oil fatty acids, benzoic acid, p-t-butylbenzoic acid, ethylbenzoic acid, isopropylbenzoic acid, toluic acid, xylylic acid, naphthenic acid and cyclohexanecarboxylic acid.

20

The overbased alkaline earth metal carboxylate/ carbonate to be used in the present invention is a compound schematically represented by the formula $\text{R}_2\text{M} \cdot \text{xMCO}_3$, as described above. However it completely differs from a mere mixture comprising a normal alkaline earth metal carboxylate and an alkaline earth metal carbonate. It is formulated into a complex through some interaction between the components thereof and characterized in that it is in the form of a homogeneous liquid while maintaining a high alkaline earth metal content.

Since the overbased alkaline earth metal carboxylate/carbonate of the present invention is in the form of a complex as described above, it can exert an excellent stabilizing effect which a mere mixture thereof can never achieve.

30

The overbased alkaline earth metal carboxylate/carbonate to be used in the present invention may be produced by a known method, for example, by reacting the above-mentioned organic carboxylic acid with a stoichiometrically excessive amount of an alkaline earth metal carbonate; by reacting an organic carboxylic acid with a stoichiometrically excessive amount of an alkaline earth metal oxide or hydroxide and then treating the resulting product with carbon dioxide; or by reacting a normal organic carboxylate of an alkaline

35

earth metal with an alkaline earth metal carbonate. The overbased alkaline earth metal carboxylate/carbonate thus produced is characterized by, for example, the metal and acid used, the content of the metal and metal ratio. The term "metal ratio" as used herein means the ratio of the total stoichiometric equivalent of the metal contained in the salt to the equivalent of the metal required for forming a neutral salt and correspond to x (as defined in the above formula) + 1.

40

The following Synthetic Examples show simple examples of the synthesis of the overbased alkaline earth metal carboxylate/carbonate to be used in the present invention.

Synthetic Example 1

45

Overbased barium oleate/carbonate

41 g of anhydrous barium hydroxide was slowly added to 300 g of methyl cellosolve and dissolved therein. Next, 11 g of oleic acid and a mixture of 10 g of nonyl phenol with 36 g of an alkylbenzene solvent were added thereto followed by stirring for approximately 30 minutes. Then the mixture was allowed to

50

react while blowing carbon dioxide into the reaction system and the temperature was slowly elevated to 150 °C while removing the water thus formed. After dehydrating and removing the methyl cellosolve under reduced pressure, overbased barium oleate/carbonate (Ba = 32 %, metal ratio = 12) was obtained as a brown viscous product.

55

Synthetic Example 2

Overbased barium octoate/carbonate

34 g of anhydrous barium hydroxide was slowly added to 480 g of methyl cellosolve and dissolved therein. Next, 6 g of octylic acid and a mixture of 20 g of nonylphenol with 34 g of an alkylbenzene solvent were added thereto followed by stirring for approximately 30 minutes. Then the mixture was allowed to react while blowing carbon dioxide into the reaction system and the temperature was slowly elevated to 150 °C while removing the water thus formed. After dehydrating and removing the methyl cellosolve under reduced pressure, overbased barious octoate/carbonate (Ba = 28 %, metal ratio = 10) was obtained as a brown viscous product.

Synthetic Example 3

10

Overbased calcium oleate/carbonate

30 g of calcium hydroxide was suspended in 300 g of methanol and the obtained suspension was allowed to react while blowing carbon dioxide into the reaction system at 15 °C. Next, 11 g of oleic acid and a mixture of 10 g of nonylphenol with 63 g of an alkylbenzene solvent were added thereto and the mixture was allowed to react by slowly elevating the temperature to 150 °C while removing the methanol and the water thus formed. After dehydrating under reduced pressure, overbased calcium oleate/carbonate (Ca = 13 %, metal ratio = 20) was obtained as a pale brown viscous product.

The overbased alkaline earth metal carboxylate/ carbonate to be used in the present invention may be easily produced by the above-mentioned methods. Alternately, various commercially available complexes may be used as such therefor. Typical examples of these commercially available complexes are LZ-2116 (overbased barium oleate/carbonate; specific gravity = 1.47, Ba = 34 %) and LZ-2117 (overbased calcium oleate/carbonate; specific gravity = 1.07, Ca = 14.2 %), each a product of Lubrizol Corp., USA.

Such an overbased alkaline earth metal carboxylate/ carbonate may be added in an amount of from 0.05 to 10 parts by weight, preferably from 0.1 to 5 parts by weight, per 100 parts by weight of the halogenated resin.

Examples of the β -diketone compound to be used in the present invention include dehydroacetic acid, acetyltetralone, benzoyltetralone, 2-benzoylcyclohexanone, dibenzoylmethane, benzoyl-4-chlorobenzoylmethane, bis(4-methylbenzoyl)methane, 4-tert-butylbenzoyl-4-methoxybenzoylmethane, benzoyl-4-isopropylbenzoylmethane, bis(4-methoxybenzoyl)methane, acetylbenzoylmethane, acetylbenzoylallylmethane, propionyl-3-methoxycarbonylbenzoylmethane, isovaleroylbenzoylmethane, octanoylbenzoylmethane, palmitoylbenzoylmethane, stearoylbenzoylmethane, dipivaloylmethane and distearoylmethane. Further, metal (for example, lithium, sodium, potassium, magnesium, calcium, barium or zinc) salts of these β -diketone compounds are similarly available in the present invention.

Such a β -diketone compound or a metal salt thereof may be used in an amount of from 0.001 to 5 parts by weight, preferably from 0.01 to 3 parts by weight, per 100 parts by weight of the halogenated resin.

Examples of the halogenated resin to be stabilized in the present invention include vinyl chloride homopolymer, copolymers of vinyl chloride with ethylene, propylene, vinyl acetate, (meth)acrylate, maleic anhydride, phenylmaleimide or cyclohexylmaleimide, polyvinylidene chloride, polyvinyl bromide, chlorinated polyethylene, graft polymer of vinyl chloride with ethylene/vinyl acetate copolymer and graft polymer of vinyl chloride with a urethane resin having an unsaturated group, each produced by a known method, for example, suspension polymerization, block polymerization or emulsion polymerization. Further, mixtures of these polymers with other thermoplastic resins are also available.

As a matter of course, the composition of the present invention may further contain metal (Li, Na, K, Ca, Ba, Mg, Sr, Zn, Cd, Sn, Cs, Al, or organic Sn) salts of carboxylic acids, organic phosphorus acids or phenols. Examples of said carboxylic acids include caproic acid, caprylic acid, pelargonic acid, 2-ethylhexylic acid, capric acid, neodecanoic acid, undecylenic acid, lauric acid, myristic acid, palmitic acid, stearic acid, isostearic acid, 12-hydroxystearic acid, chlorostearic acid, 12-ketostearic acid, phenylstearic acid, ricinolic acid, linoleic acid, linolenic acid, oleic acid, arachic acid, behenic acid, erucic acid, brassidic acid and similar acids, mixtures of the above-cited acids with natural acids such as tallow fatty acids, coconut oil fatty acids, tung oil fatty acids, soybean oil fatty acids and cottonseed oil fatty acids, benzoic acid, p-t-butylbenzoic acid, ethylbenzoic acid, isopropylbenzoic acid, toluic acid, xylylic acid, salicylic acid, 5-t-octylsalicylic acid, naphthenic acid and cyclohexanecarboxylic acid. Examples of said organic phosphorus acids include mono- or dioctylphosphoric acid, mono- or di-dodecylphosphoric acid, mono- or dioctadecylphosphoric acid, mono- or di-(nonylphenyl)phosphoric acid, nonyl phosphonate and stearyl phosphonate. Examples of said phenols include phenol, cresol, ethylphenol, cyclohexylphenol, nonylphenol and dodecylphenol.

Such a metal salt may be used in an amount of preferably from 0.05 to 10 parts by weight per 100

parts by weight of the halogenated resin.

The composition of the present invention may further contain various additives commonly used as additives for halogenated resins, for example, organic phosphite compounds and/or epoxy compounds and plasticizer.

- 5 Examples of said organic phosphite compounds include triphenyl phosphite, tris(nonylphenyl) phosphite, tris(mono- and di-mixed nonylphenyl) phosphite, diphenyl decyl phosphite, diphenyl tridecyl phosphite, phenyl dioctyl phosphite, phenyl diisodecyl phosphite, phenyl di(tridecyl) phosphite, tridecyl phosphite, dibutyl acid phosphite, dilauryl acid phosphite, diphenyl acid phosphite, di(nonylphenyl) acid phosphite, distearyl pentaerythritol diphosphite, di(tridecyl) pentaerythritol diphosphite, phenyl-4,4'-10 isopropylidenediphenol pentaerythritol diphosphite, tetra (C_{12-15} mixed alkyl)-4,4'-isopropylidenediphenyl diphosphite and hydrogenated 4,4'-isopropylidenediphenol polyphosphite.

Such an organic phosphite may be used in an amount of preferably from 0.01 to 5 parts by weight per 100 parts by weight of the halogenated resin.

- 15 Examples of said epoxy compounds include epoxidized animal and vegetable oils such as epoxy soybean oil, epoxy linseed oil, epoxydized tung oil, epoxidized fish oil, epoxydized beef tallow oil and epoxidized safflower oil, methyl epoxystearate, butyl epoxystearate, 2-ethylhexyl epoxystearate, stearyl epoxystearate, epoxidized polybutadiene, tris(epoxypropyl) isocyanurate, epoxidized tall oil fatty acid ester, epoxidized linseed oil fatty acid ester, bisphenol A diglycidyl ether, vinylcyclohexene diepoxide, dicyclohexene diepoxide and 3,4-epoxycyclohexylmethyl epoxycyclohexanecarboxylate.

- 20 Examples of said plasticizer include phthalate plasticizers such as diheptyl phthalate, dioctyl phthalate and diisononyl phthalate, adipate plasticizers such as diethyl adipate, diisononyl adipate and di-(butyldiglycol) adipate, phosphate plasticizers, polyester plasticizers, chlorinated paraffin plasticizers and trimellitate plasticizers.

- 25 In addition, the composition of the present invention may further contain various additives commonly used in chlorinated vinyl resins, for example, crosslinking agent, filler, foaming agent, antistatic, antifogging agent, plate-out inhibitor, surface treatment, lubricant, flame-retardant, fluorescent agent, mildewproofing agent, bactericide, sequestering agent, mold releasing agent, pigment, processing aid, antioxidant, light stabilizer, seolites, powdery silicate (silica), perchlorates such as sodium perchlorate, magnesium perchlorate and barium perchlorate and hydrotalcites such as synthetic hydrotalcite, zinc-modified hydrotalcite and perchloric acid-treated (zinc-modified) hydrotalcite.

The composition of the present invention may be used regardless of the processing method of the halogenated resin. For example, it is highly applicable to calendering, rolling, extrusion molding, pressure molding, paste processing and powder molding.

- 30 To further illustrate the present invention, and not by way of limitation, the following Examples will be given.

Example 1

- 35 The composition as specified below was kneaded on a roll at 180 °C for 7 minutes and then pressed at 40 170 °C for 5 minutes to thereby form a sheet of 0.5 mm in thickness.

Test pieces were cut from this sheet and then subjected to the following tests.

For comparison, pieces prepared by the same method as the above-mentioned one except that the LZ-2116 was replaced by LZ-2106 (a product of Lubrizol Corp., USA; overbased barium nonylphenate/carbonate) were similarly tested.

- 45 Table 1 shows the results.

(1) Initial coloration:

- 40 The yellowness of a test piece was measured.

(2) Heat stability:

- 45 A test piece was heated in a Geer oven at 190 °C and the time required for blackening was measured.

50 (3) Adhesiveness:

A vinyl acetate adhesive was applied to the aluminum face of a polyester film metallized with aluminum. Next, a test piece was adhered thereto and preserved at 70 °C and a relative humidity of 98 % for several

days. Then the adhesion of the test piece to the aluminum foil was observed. The result was evaluated in ten grades [1 (no peeling) < _____ > 10 (easy peeling of the whole test piece)].

5	(Composition)	part by weight
	polyvinyl chloride resin	95
10	(Zeon 103 EP-8, a product of Nippon Zeon)	
	acrylic impact resistance modifier	5
	(Kaneace B-31, a product of Kanegafuchi	
15	Chemical Industry)	
	dioctyl phthalate	8
20	octyl epoxystearate	4
	zinc stearate	0.2
	barium stearate	0.5
25	zinc benzoate	0.2
	phenyl di(tridecyl) phosphite	1.5
30	LZ-2116* ¹	0.5
	β -diketone compound	0.4

35 *1: a product of Lubrizol Corp., USA, overbased
 barium oleate/carbonate.

40

45

50

55

5
10
15
20
25
30
35
40
45
50

Table 1

No.	β -Diketone compound	Initial coloration	Heat stability (min)	Adhesiveness		
				5 days	10 days	20 days
Comp. Ex. LZ-2106 compound system						
1 - 1	none	26.8	70	4	7	10
1 - 2	dibenzoylmethane	22.7	85	4	7	10
1 - 3	stearylbenzoylmethane	23.5	90	4	7	10
Comp. Ex. LZ-2116 compound system						
1 - 4	none	26.4	80	3	6	10
Ex.						
1 - 1	dibenzoylmethane	17.6	120	1	1	3
1 - 2	stearylbenzoylmethane	17.5	120	1	1	3
1 - 3	3-methoxycarbonyl- benzoylpropionylmethane	17.2	120	1	3	5
1 - 4	dehydroacetic acid	18.9	115	1	3	6
1 - 5	zinc dehydroacetate	18.4	110	1	2	6

55 Example 2

By using the composition as specified below, test pieces were prepared in the same manner as the one described in Example 1. Then the heat stability and adhesiveness of the obtained test piece were tested in

the same manner as the one described in Example 1. In the case of the initial coloration test, the whiteness of the test piece was measured.

	(Composition)	part by weight
5	polyvinyl chloride resin (Zeon 103 EP-8, a product of Nippon Zeon)	95
10	acrylic impact resistance modifier (Kaneace B-31, a product of Kanegafuchi Chemical Industry)	5
15	dioctyl phthalate	8
20	octyl epoxystearate	4
25	zinc stearate	0.2
	barium stearate	0.5
30	zinc benzoate	0.2
	tetra(C ₁₂₋₁₅ alkyl)bisphenol A diphosphite	1.5
	calcium carbonate	5
35	stearoylbenzoylmethane	0.4
	sample compound	0.5
40		
45		
50		
55		

Table 2

No.	Sample compound	Initial coloration	Heat stability (min)	Adhesiveness		
				5 days	10 days	20 days
Comp. Ex.						
2 - 1	barium oleate	84.5	90	4	8	10
2 - 2	basic barium oleate	83.4	85	4	6	10
2 - 3	LZ-2116	87.7	100	4	7	10
2 - 4	barium oleate (0.1) barium carbonate (0.4)	82.6	80	5	9	10
2 - 5	calcium oleate	83.3	75	4	8	10
Ex.						
2 - 1	LZ-2116	92.6	120	1	1	3
2 - 2	LZ-2116	91.8	110	1	1	3
2 - 3	product of Syn. Ex. 1	91.3	120	1	2	4
2 - 4	product of Syn. Ex. 2	91.5	115	1	2	5
2 - 5	product of Syn. Ex. 3	91.1	110	1	2	5

55 Example 3

By using the composition as specified below, test pieces were prepared in the same manner as the one described in Example 1. Then the heat stability of the test piece was measured in a Geer oven at 190 °C.

In order to evaluate the dynamic heat stability, the time required for the sticking of the compound to the roll (i.e., sticking time) was measured and the extent of coloration (mill colorability) was observed with the lapse of time. The result was evaluated in ten grades [1 (no coloration) → 10 (remarkable coloration)].

- 5 For comparison, the LZ-2116 was replaced with LZ-2106 and the obtained pieces were similarly tested. Table 3 shows the results.

	(Composition)	part by weight
10	polyvinyl chloride resin (Zeon 103 EP-8, a product of Nippon Zeon)	95
15	acrylic impact resistance modifier (Kaneace B-31, a product of Kanegafuchi Chemical Industry)	5
20	dioctyl phthalate	8
	epoxidized soybean oil	4
25		
	zinc stearate	0.2
30	barium stearate	0.5
	zinc benzoate	0.2
35	diphenyl tridecyl phosphite	1.0
	LZ-2116	0.6
	β-diketone compound	0.4
40		
45		
50		

5
10
15
20
25
30
35
40
45
50

Table 3

No.	β -diketone compound	Heat stability (min)	Sticking time (min)	Melt colorability		
				5 days	10 days	20 days
Comp. Ex. LZ-2106 compound system						
3 - 1	none	60	23.5	3	6	9
3 - 2	dibenzoylmethane	75	26.5	1	3	6
3 - 3	stearoylbenzoylmethane	70	26.0	1	4	7
Comp. Ex. LZ-2116 compound system						
3 - 4	none	65	25.5	3	5	9
Ex.						
3 - 1	dibenzoylmethane	105	31.5	1	2	3
3 - 2	zinc dibenzoylmethane	100	30.0	1	2	3
3 - 3	stearoylbenzoylmethane	105	31.0	1	2	3
3 - 4	isopivaloylbenzoylmethane	100	30.5	1	2	4
3 - 5	dehydroacetic acid	95	30.0	1	2	4

55 Example 4

The composition as specified below was kneaded on a roll at 160 °C for 5 minutes and then pressed at 160 °C for 5 minutes to thereby form a sheet of 0.5 mm in thickness.

Test pieces were cut from this sheet and then subjected to the following tests. Table 4 shows the results.

(1) Heat stability:

5 A test piece was heated in a Geer oven at 190 °C and the time required for blackening was measured.

(2) Thermal coloration:

10 The coloration of a test piece heated at 190 °C for 60 minutes was evaluated in ten grades [1 (no coloration) < _____ > 10 (remarkable coloration)].

(3) Heat aging characteristics:

15 A test piece was heated in a Geer oven at 110 °C and the time required for blackening was measured.

	(Composition)	part by weight
20	polyvinyl chloride resin (Zeon 103 EP, a product of Nippon Zeon)	100
25	ABS resin (Blendex, a product of Ube Cycon, Ltd.)	5
30	di(C ₉₋₁₁ alkyl) phthalate	70
35	epoxyidized soybean oil	2
40	antimony trioxide	2
	zinc stearate	0.1
	barium stearate	0.2
45	zinc benzoate	0.2
	barium perchlorate	0.1
50	phenyl di(tridecyl) phosphite	0.6
	LZ-2116	0.2
	β-diketone compound	0.1

Table 4

No.	β -Diketone compound	Heat stability (min)	Thermal coloration	Heat aging characteristics (day)
Comp. Ex.	LZ-2106 compound system			
4 - 1	none	75	9	16
4 - 2	dibenzoylmethane	90	6	19
4 - 3	stearoylbenzoylmethane	90	6	18
Comp. Ex.	LZ-2116 compound system			
4 - 4	none	80	8	17
Ex.				
4 - 1	dibenzoylmethane	120	3	28
4 - 2	zinc dibenzoylmethane	110	3	26
4 - 3	stearoylbenzoylmethane	115	3	26
4 - 4	isopivaloylbenzoylmethane	115	4	25
4 - 5	dehydroacetic acid	105	4	25

55 Example 5

The composition as specified below was kneaded on a roll at 180 °C for 7 minutes and then pressed at 170 °C for 5 minutes to thereby form a sheet of 0.5 mm in thickness.

Test pieces were cut from this sheet and then subjected to the following tests.
Table 5 shows the results.

(1) Heat stability:

5 A test piece was heated in a Geer oven at 190 °C and the time required for blackening was measured.

(2) Light resistance:

10 The colorations of test pieces irradiated with light in a fadeometer at a black panel temperature of 83 °C for 10 and 25 days were evaluated in ten grades [1 (no coloration < _____) 10 (remarkable coloration)].

(3) Heat aging characteristics:

15 A test piece was heated in a Geer oven at 110 °C for 2 weeks and the coloration thereof was evaluated in ten grades [1 (no coloration) < _____) 10 (remarkable coloration)].

	(Composition)	part by weight
20	polyvinyl chloride resin	100
	(TK-1300; a product of The Shin-Etsu	
25	Chemical)	
	di(C ₉₋₁₁ alkyl) phthalate	80
30	chlorinated paraffin	10
	calcium carbonate	15
	epoxidized soybean oil	4
35	zinc stearate	0.4
	barium stearate	0.5
40	phenyl di(tridecyl) phosphite	0.5
	dibenzoylmethane	0.05
	sample compound	0.4

45

50

55

Table 5

No.	Sample compound	Heat stability (min)	Light resistance		Heat aging characteristics (day)
			10 days	25 days	
Comp. Ex. 5 - 1	barium oleate	75	7	9	9
5 - 2	basic barium oleate	80	4	9	8
5 - 3	LZ-2106	85	4	7	8
5 - 4	barium oleate (0.1) barium carbonate (0.4)	65	7	10	10
5 - 5	calcium oleate	70	6	9	9
Ex. 5 - 1	LZ-2116	105	2	3	4
5 - 2	LZ-2117	95	2	3	5
5 - 3	product of Syn. Ex. 1	100	2	3	4
5 - 4	product of Syn. Ex. 2	100	2	3	5
5 - 5	product of Syn. Ex. 3	90	3	4	5

The results of the above Examples indicate the following facts.

When an overbased phenolate/carbonate, which is an overbased complex other than those to be used in the present invention, was used, the obtained composition showed somewhat improved but yet unsatisfactory heat stability and coloration. In this case, furthermore, the adhesiveness of the obtained product, in particular, to a different kind of material was seriously insufficient. When a basic metal

carboxylate was used or a metal carboxylate was used together with a metal carbonate, scarcely any improvement was achieved as compared with a case wherein a conventional metal carboxylate was used. In this case, further, the adhesiveness was seriously poor.

In contrast, the composition of the present invention comprising an overbased metal carboxylate/carbonate together with a β -diketone compound showed not only extremely improved heat stability and coloration but also an excellent adhesiveness to a different kind of material, which clearly shows that the effects of the present invention are remarkably excellent.

Claims

10

1. A stabilized halogenated resin composition which comprises 100 parts by weight of a halogenated resin and, added thereto, (a) 0.05 to 10 parts by weight of at least one overbased alkaline earth metal carboxylate/carbonate and (b) 0.001 to 5 parts by weight of at least one β -diketone compound or a metal salt thereof.

15

2. A halogenated resin composition as claimed in Claim 1, wherein said overbased alkaline earth metal carboxylate/carbonate is an overbased calcium or barium carboxylate/carbonate.

20

3. A halogenated resin composition as claimed in Claim 2, wherein said overbased barium carboxylate/carbonate is overbased barium oleate/carbonate.

4. A halogenated resin composition as claimed in Claim 2, wherein said overbased calcium carboxylate/carbonate is overbased calcium oleate/carbonate.

25

5. A halogenated resin composition as claimed in Claim 1, wherein said β -diketone compound is dibenzoylmethane.

6. A halogenated resin composition as claimed in Claim 1, wherein said β -diketone compound is stearoylbenzoylmethane.

30

7. A halogenated resin composition as claimed in Claim 1, wherein said β -diketone compound is dehydroacetic acid.

8. A halogenated resin composition as claimed in Claim 1, which further comprises a zinc carboxylate as an additional stabilizer.

9. A halogenated resin composition as claimed in Claim 1, wherein said halogenated resin is a polyvinyl chloride resin.

40

45

50

55



European
Patent Office

EUROPEAN SEARCH
REPORT

Application Number

EP 91 10 2550

DOCUMENTS CONSIDERED TO BE RELEVANT					
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)		
P,X	EP-A-0 394 547 (AKISHIMA CHEMICAL INDUSTRIES CO., LTD.) * claims; page 4, lines 42-44 * -----	1-9	C 08 K 13/02 C 08 L 57/08 // (C 08 K 13/02 C 08 K C 08 K 3:26 C 08 K 5:07 C 08 K C 08 K 5:09)		
D,A	FR-A-2 423 510 (S.A. ARGUS CHEMICAL NV.) * claims * -----	1-9			
The present search report has been drawn up for all claims			TECHNICAL FIELDS SEARCHED (Int. Cl.5)		
			C 08 K C 08 L		
Place of search Date of completion of search Examiner					
The Hague	21 June 91	DE LOS ARCOS Y VELAZ			
CATEGORY OF CITED DOCUMENTS					
X: particularly relevant if taken alone					
Y: particularly relevant if combined with another document of the same category					
A: technological background					
O: non-written disclosure					
P: intermediate document					
T: theory or principle underlying the invention					
E: earlier patent document, but published on, or after the filing date					
D: document cited in the application					
L: document cited for other reasons					
S: member of the same patent family, corresponding document					